7 Appendix: regulatory information

7.1 Medical devices of other manufacturers

Please note that this manual may also contain information about medical devices that are NOT legally manufactured by Siemens. These medical devices are either only distributed by Siemens, or only mentioned for additional information.

In the following table you will find information about the medical device, the corresponding legal manufacturer and, if applicable, the authorized representative.

Product and CE	Legal manufacturer
Respiration cushion, respiratory belt CE	Pi-Products GmbH Heinrich-Hertz-Straße 8a 92224 Amberg, Germany www.pi-products.de
Application cushion for PMU CE	Polyform GmbH & Co. KG Kunststofftechnik Braasstraße 15 DE-31737 Rinteln, Germany www.polyform.de
Patient table CE	TRUMPF Medizin Systeme GmbH + Co.KG Carl-Zeiss-Str. 7-9 07318 Saalfeld, Germany www.de.trumpf.com

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7.2 CE for Physiological Measurement Unit (PMU)

The PMU (PPU, PERU) bears a CE marking in accordance with the provisions of regulation 1999/5/EC of March 9, 1999 for radio and telecommunications terminal equipment.

7.3 FCC and IC for Physiological Measurement Unit

Changes or modifications not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment.

This device complies with Industry Canada licence-exempt RSS standard(s) and part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) this device may not cause interference, and (2) this device must accept any interference, including interference that may cause undesired operation of the device.

Le présent appareil est conforme aux CNR d'Industrie Canada applicables aux appareils radio exempts de licence. L'exploitation est autorisée aux deux conditions suivantes :
(1) l'appareil ne doit pas produire de brouillage, et
(2) l'appareil doit accepter tout brouillage radioélectrique subi, même si le brouillage est susceptible d'en compromettre le fonctionnement.

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